Notice of Allowability	Application No.	Applicant(s)
	09/749,410	NERIISHI ET AL.
	Examiner	Art Unit
	Carla Myers	1634
The MAILING DATE of this communication appe All claims being allowable, PROSECUTION ON THE MERITS IS herewith (or previously mailed), a Notice of Allowance (PTOL-85) NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT R of the Office or upon petition by the applicant. See 37 CFR 1.313	(OR REMAINS) CLOSED in this or other appropriate communical IGHTS. This application is subject	application. If not included tion will be mailed in due course. THIS
1. This communication is responsive to		
2. The allowed claim(s) is/are 1-3 and 8.		
3. The drawings filed on 28 December 2000 are accepted by the Examiner.		
4.		
Attachment(s)  1. Notice of References Cited (PTO-892)  2. Notice of Draftperson's Patent Drawing Review (PTO-948)  3. Information Disclosure Statements (PTO-1449 or PTO/SB/06 Paper No./Mail Date  4. Examiner's Comment Regarding Requirement for Deposit of Biological Material	6. ☐ Interview Summa Paper No./Mail D 3), 7. ⊠ Examiner's Amen	Date .

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The Notice of Withdrawal from Issue dated May 5, 2004 was mailed in error.

The Office sincerely apologizes for any inconvenience that this error may have caused.

The examiner's amendment set forth in the previous Notice of Allowance of April 21,

2004 is repeated below.

## **EXAMINER'S AMENDMENT**

An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Susan Dadio on April 14, 2004.

The application has been amended as follows:

1. A process for detecting a complementary DNA fragment which comprises the steps of:

bringing a liquid phase comprising single-stranded sample DNA fragments having a radioactive label [in a liquid phase] into contact with a DNA micro-array having a support and at least two defined areas in each of which a group of probe compounds selected from the group consisting of DNA molecules, DNA fragments, synthesized oligonucleotides, synthesized polynucleotides, and PNA (peptide nucleic acid) are fixed under such conditions that a group of the probe compounds fixed in one area differs from a group of the probe compounds fixed in another area, so that DNA fragments

complementary to a group of the probe compounds are fixed [by hybridization to the area in which the last- mentioned group is fixed] to an area of the micro-array by hybridization of complementary DNA fragments to the probe compounds;

removing unfixed sample DNA fragments from the DNA micro-array; keeping the DNA micro-array in contact with a radiation image storage panel containing a stimulable phosphor via a spacer sheet [having openings in areas corresponding to the areas on which groups of the probe compounds are fixed,] intervening between the DNA micro-array and the radiation image storage panel, said spacer sheet being in direct contact with the micro-array and having openings aligned with the areas of the micro-array to which the probe compounds are fixed, so that the radiation image storage panel can absorb and store radiation energy [of the radioactive label coming from the fixed] transmitted by the radioactive label of the fixed complementary DNA fragments through the openings in said spacer sheet;

irradiating the radiation image storage panel with a stimulating light, so that the image storage panel releases a stimulated emission from the area in which the radiation energy is stored;

detecting the stimulated emission photoelectrically to obtain a series of electric signals; and

processing the electric signals to locate the area in which the complementary DNA fragments are fixed.

8. A process for detecting a complementary DNA fragment which comprises the steps of:

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bringing a liquid phase comprising single-stranded sample DNA fragments having a radioactive label [in a liquid phase] into contact with a DNA micro-array having a support and at least two defined areas in each of which a group of probe compounds selected from the group consisting of DNA molecules, DNA fragments, synthesized oligonucleotides, synthesized polynucleotides, and PNA (peptide nucleic acid), are fixed under such conditions that a group of the probe compounds fixed in one area differs from a group of the probe compounds fixed in another area, so that DNA fragments complementary to a group of the probe compounds are fixed [by hybridization to the area in which the last- mentioned group is fixed] to an area of the micro-array by hybridization of complementary DNA fragments to the probe compounds;

removing unfixed sample DNA fragments from the DNA micro-array;

keeping the DNA micro-array in contact with a radiation image storage panel containing a stimulable phosphor via a spacer sheet [having openings in areas corresponding to the areas on which groups of the probe compounds are fixed,] intervening between the DNA micro-array and the radiation image storage panel, said spacer sheet being in direct contact with the micro-array and having openings aligned with the areas of the micro-array to which the probe compounds are fixed, so that the radiation image storage panel can absorb and store radiation energy [of the radioactive label coming from the fixed] transmitted by the radioactive label of the fixed complementary DNA fragments through the openings in said spacer sheet;

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irradiating the radiation image storage panel with a stimulating light, so that the image storage panel releases a stimulated emission from the area in which the radiation energy is stored;

detecting the stimulated emission photoelectrically to obtain a series of electric signals; and

processing the electric signals to locate the area in which the complementary DNA fragments are fixed,

wherein said spacer sheet has a thickness in the range of 10 to 300 µm and is made of a non radiation-transmitting material [is] selected from the group consisting of aluminum, brass, stainless steel, polyethylene terephthalate and polyethylene naphthalate.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Carla Myers whose telephone number is (571) 272-0747. The examiner can normally be reached on Monday-Thursday from 6:30 AM-5:00 PM. A message may be left on the examiner's voice mail service. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion, can be reached on (571)-272-0782.

Papers related to this application may be faxed to Group 1634 via the PTO Fax Center using the fax number (703)-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Carla Myers May 19, 2004

CARLA J. MYERS
PRIMARY EXAMINER